

=> s us6071966/pn  
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4 219811-13-9/BI  
4 219811-14-0/BI  
3 219811-15-1/BI  
3 219811-16-2/BI  
4 219811-17-3/BI

*Q2704*  
*Instant*  
*Compounds*

4 219811-18-4/BI  
 3 219811-19-5/BI  
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*all below having the instant compound*

L2 ANSWER 1 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2004:372886 CAPLUS  
 DOCUMENT NUMBER: 140:368722  
 TITLE: Combination therapy using 1-aminocyclohexane derivatives and acetylcholinesterase inhibitors for treatment of dementia  
 INVENTOR(S): Moebius, Hans-Joerg  
 PATENT ASSIGNEE(S): Germany  
 SOURCE: U.S. Pat. Appl. Publ., 46 pp.  
 CODEN: USXXCO  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 2  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004087658	A1	20040506	US 2003-691895	20031023
PRIORITY APPLN. INFO.:			US 2002-420918P	P 20021024

L2 ANSWER 2 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2004:368913 CAPLUS  
 DOCUMENT NUMBER: 140:395498  
 TITLE: Preparation and combination therapy of cyclohexanamines and acetylcholinesterase inhibitors for treatment of dementia  
 INVENTOR(S): Moebius, Hans-Joerg  
 PATENT ASSIGNEE(S): Merz Pharma G.m.b.H. & Co. K.-G.a.A., Germany;  
 Marsden, John Christopher  
 SOURCE: PCT Int. Appl., 113 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 2  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004037234	A2	20040506	WO 2003-GB4549	20031023
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,				

CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC,  
NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,  
GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.:

US 2002-420918P P 20021024

L2 ANSWER 3 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2004:353140 CAPLUS

DOCUMENT NUMBER: 140:380634

TITLE: Compositions of cyclooxygenase-2 selective inhibitors  
and NMDA receptor antagonists for the treatment or  
prevention of neuropathic pain

INVENTOR(S): Cheung, Raymond Y.

PATENT ASSIGNEE(S): Pharmacia Corporation, USA

SOURCE: U.S. Pat. Appl. Publ., 51 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004082543	A1	20040429	US 2002-282660	20021029
WO 2004039371	A2	20040513	WO 2003-US33089	20031017
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.:

US 2002-282660 A 20021029

L2 ANSWER 4 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2004:270008 CAPLUS

DOCUMENT NUMBER: 140:297535

TITLE: Methods of treating age-associated memory impairment,  
mild cognitive impairment, and dementias with cell  
cycle inhibitors

INVENTOR(S): Reisberg, Barry

PATENT ASSIGNEE(S): New York University, USA

SOURCE: PCT Int. Appl., 40 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004026246	A2	20040401	WO 2003-US29403	20030917
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC,			

NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,  
GW, ML, MR, NE, SN, TD, TG  
PRIORITY APPLN. INFO.: US 2002-411282P P 20020917

L2 ANSWER 5 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2004:80486 CAPLUS  
DOCUMENT NUMBER: 140:139523  
TITLE: NMDA receptor antagonists and their use in inhibiting  
abnormal hyperphosphorylation of protein Tau  
INVENTOR(S): Iqbal, Khalid; Grundke-Iqbal, Inge  
PATENT ASSIGNEE(S): USA  
SOURCE: PCT Int. Appl., 97 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004009062	A2	20040129	WO 2003-US22362	20030717
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
US 2004019118	A1	20040129	US 2003-622163	20030717
PRIORITY APPLN. INFO.:			US 2002-397434P	P 20020719
OTHER SOURCE(S):	MARPAT 140:139523			

L2 ANSWER 6 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2003:777580 CAPLUS  
DOCUMENT NUMBER: 139:292154  
TITLE: Preparation of azabicyclic derivatives of  
aminocyclohexane as NMDA, 5HT3, and neuronal nicotinic  
receptor antagonists  
INVENTOR(S): Parsons, Christopher Graham Raphael; Henrich, Markus;  
Danyasz, Wojciech; Kalvinsh, Ivars; Kauss, Valerjans;  
Jirgensons, Aigars; Gold, Markus; Vanejevs, Maksims  
PATENT ASSIGNEE(S): Merz Pharma G.m.b.H. & Co. K.-G.a.A., Germany  
SOURCE: PCT Int. Appl., 91 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003080046	A1	20031002	WO 2003-GB1236	20030321
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,			

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NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,  
GW, ML, MR, NE, SN, TD, TG  
X US 2004034055 A1 20040219 US 2003-394670 20030321  
PRIORITY APPLN. INFO.: US 2002-366386P P 20020321  
OTHER SOURCE(S): MARPAT 139:292154  
REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 7 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2003:376808 CAPLUS  
DOCUMENT NUMBER: 138:379247  
TITLE: Unsaturated 1-amino-alkylcyclohexane NMDA, 5HT3 and  
neuronal nicotinic receptor antagonists  
INVENTOR(S): Parsons, Christopher Graham Raphael; Henrich, Markus;  
Dansyz, Wojciech; Kalvinsh, Ivars; Kauss, Valerjans;  
Jirgensons, Aigars  
PATENT ASSIGNEE(S): Merz Pharma Gmbh & Co. Kgaa, Germany; Gold, Markus  
SOURCE: PCT Int. Appl., 104 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003040084	A1	20030515	WO 2002-GB5038	20021107
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			

X US 2003166634 A1 20030904 US 2002-288819 20021106  
PRIORITY APPLN. INFO.: US 2001-350974P P 20011107  
US 2001-337858P P 20011108  
OTHER SOURCE(S): MARPAT 138:379247  
REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 8 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2003:249800 CAPLUS  
DOCUMENT NUMBER: 139:173649  
TITLE: Are neuronal nicotinic receptors a target for  
antiepileptic drug development? Studies in different  
seizure models in mice and rats  
AUTHOR(S): Loscher, Wolfgang; Potschka, Heidrun; Wlaz, Piotr;  
Danyasz, Wojciech; Parsons, Christopher G.  
CORPORATE SOURCE: Toxicology and Pharmacy, Department of Pharmacology,  
School of Veterinary Medicine, Hannover, 30559,  
Germany  
SOURCE: European Journal of Pharmacology (2003), 466(1-2),  
99-111  
CODEN: EJPHAZ; ISSN: 0014-2999  
PUBLISHER: Elsevier Science B.V.  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
REFERENCE COUNT: 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 9 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2002:725218 CAPLUS  
 DOCUMENT NUMBER: 138:297293  
 TITLE: NMDA receptor antagonists to characterize rat renal  
 organic cation transporter function  
 AUTHOR(S): Fourie, Jeanne; Escobar, Miguel R.; Sitar, Daniel S.  
 CORPORATE SOURCE: Department of Pharmacology and Therapeutics,  
 University of Manitoba, Winnipeg, MB, R3E 0W3, Can.  
 SOURCE: European Journal of Pharmacology (2002), 452(1), 1-10  
 CODEN: EJPHAZ; ISSN: 0014-2999  
 PUBLISHER: Elsevier Science B.V.  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS  
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 10 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2002:546227 CAPLUS  
 DOCUMENT NUMBER: 138:180507  
 TITLE: Synergistic effect of uncompetitive NMDA receptor  
 antagonists and antidepressant drugs in the forced  
 swimming test in rats  
 AUTHOR(S): Rogoz, Zofia; Skuza, Grazyna; Maj, Jerzy; Danysz,  
 Wojciech  
 CORPORATE SOURCE: Institute of Pharmacology, Polish Academy of Sciences,  
 Krakow, PL 31-343, Pol.  
 SOURCE: Neuropharmacology (2002), 42(8), 1024-1030  
 CODEN: NEPHBW; ISSN: 0028-3908  
 PUBLISHER: Elsevier Science Ltd.  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 REFERENCE COUNT: 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS  
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 11 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2002:329206 CAPLUS  
 DOCUMENT NUMBER: 137:241556  
 TITLE: Amino-alkyl-cyclohexanes as a novel class of  
 uncompetitive NMDA receptor antagonists  
 AUTHOR(S): Danysz, W.; Parsons, C. G.; Jirgensons, A.; Kauss, V.;  
 Tillner, J.  
 CORPORATE SOURCE: Merz+Co., Frankfurt am Main, 60318, Germany  
 SOURCE: Current Pharmaceutical Design (2002), 8(10), 835-843  
 CODEN: CPDEFP; ISSN: 1381-6128  
 PUBLISHER: Bentham Science Publishers  
 DOCUMENT TYPE: Journal; General Review  
 LANGUAGE: English  
 REFERENCE COUNT: 43 THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS  
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 12 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2002:240553 CAPLUS  
 DOCUMENT NUMBER: 136:268173  
 TITLE: 1-Aminoalkylcyclohexanes as trypanocidal agents  
 INVENTOR(S): Kelly, John M.; Kalvinsh, Ivars; Kauss, Valerjans;  
 Jirgensons, Aigars; Gold, Markus  
 PATENT ASSIGNEE(S): Merz & Co. G.m.b.H. & Co., Germany  
 SOURCE: PCT Int. Appl., 32 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002024186	A1	20020328	WO 2001-EP10731	20010914
W: AU, CA, CN, CZ, CZ, GE, HU, IL, JP, KR, MX, NO, PL, UA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR				
US 6602862	B1	20030805	US 2000-664629	20000919
AU 2001087740	A5	20020402	AU 2001-87740	20010914
EP 1318800	A1	20030618	EP 2001-967349	20010914
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY, TR				
JP 2004509144	T2	20040325	JP 2002-528257	20010914
ZA 2001007649	A	20020514	ZA 2001-7649	20010917
NO 2003001239	A	20030318	NO 2003-1239	20030318
PRIORITY APPLN. INFO.:			US 2000-664629	A 20000919
			WO 2001-EP10731	W 20010914
OTHER SOURCE(S):		MARPAT 136:268173		
REFERENCE COUNT:		9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT		

L2 ANSWER 13 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:935560 CAPLUS

DOCUMENT NUMBER: 136:48466

TITLE: 1-Aminoalkylcyclohexanes as 5-HT<sub>3</sub> and neuronal nicotinic receptor antagonists, preparation, pharmaceutical compositions, and therapeutic use thereof

INVENTOR(S): Parsons, Christopher Graham Raphael; Danysz, Wojciech; Gold, Markus; Kalvinsh, Ivars; Kauss, Valerjans; Jirgensons, Aigars

PATENT ASSIGNEE(S): Merz & Co. G.m.b.H. & Co., Germany

SOURCE: PCT Int. Appl., 40 pp.  
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001098253	A2	20011227	WO 2001-EP6964	20010619
W: AU, CA, CN, CZ, CZ, FI, FI, GE, HU, IL, JP, KR, MX, NO, PL, UA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR				
ZA 2001004187	A	20021122	ZA 2001-4187	20010522
EP 1303477	A2	20030423	EP 2001-960342	20010619
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY, TR				
JP 2004501130	T2	20040115	JP 2002-504209	20010619
NO 2002006103	A	20030219	NO 2002-6103	20021219
PRIORITY APPLN. INFO.:			US 2000-597102	A 20000620
			WO 2001-EP6964	W 20010619
OTHER SOURCE(S):		MARPAT 136:48466		

L2 ANSWER 14 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:429280 CAPLUS

DOCUMENT NUMBER: 135:251854

TITLE: The N-methyl-d-aspartate receptor channel blockers memantine, MRZ 2/579 and other amino-alkyl-cyclohexanes antagonize 5-HT<sub>3</sub> receptor currents in



cultured HEK-293 and N1E-115 cell systems in a  
 non-competitive manner  
 AUTHOR(S): Rammes, G.; Rupprecht, R.; Ferrari, U.;  
 Zieglgansberger, W.; Parsons, C. G.  
 CORPORATE SOURCE: Max-Planck-Institute of Psychiatry, Munchen, D-80804,  
 Germany  
 SOURCE: Neuroscience Letters (2001), 306(1-2), 81-84  
 CODEN: NELED5; ISSN: 0304-3940  
 PUBLISHER: Elsevier Science Ireland Ltd.  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS  
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 15 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2001:338511 CAPLUS  
 DOCUMENT NUMBER: 134:340433  
 TITLE: Preparation of N-(alkylcyclohexyl)azacycloalkanes as  
 anticonvulsants  
 INVENTOR(S): Gold, Markus; Danysz, Wojciech; Parsons, Christopher  
 Graham Raphael; Kalvinsh, Ivars; Kauss, Valerjans;  
 Jirgensons, Aigars  
 PATENT ASSIGNEE(S): Merz & Co. GmbH & Co., Germany  
 SOURCE: PCT Int. Appl., 30 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001032640	A1	20010510	WO 1999-EP8317	19991101
W:	AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
EP 1228052	A1	20020807	EP 1999-974146	19991101
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL			
JP 2003513083	T2	20030408	JP 2001-534791	19991101
NO 2002002044	A	20020430	NO 2002-2044	20020430
PRIORITY APPLN. INFO.:			WO 1999-EP8317	W 19991101
OTHER SOURCE(S):	MARPAT 134:340433			
REFERENCE COUNT:	3	THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT		

L2 ANSWER 16 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
 ACCESSION NUMBER: 2001:319290 CAPLUS  
 DOCUMENT NUMBER: 135:101928  
 TITLE: In vitro and in vivo activities of aminoadamantane and  
 aminoalkylcyclohexane derivatives against Trypanosoma  
 brucei  
 AUTHOR(S): Kelly, John M.; Quack, Guenter; Miles, Michael M.  
 CORPORATE SOURCE: Department of Infectious and Tropical Diseases, London  
 School of Hygiene and Tropical Medicine, London, WC1E  
 7HT, UK  
 SOURCE: Antimicrobial Agents and Chemotherapy (2001), 45(5),  
 1360-1366

CODEN: AMACCQ; ISSN: 0066-4804  
PUBLISHER: American Society for Microbiology  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 17 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2000:535984 CAPLUS  
DOCUMENT NUMBER: 133:281550  
TITLE: Synthesis and structure-affinity relationships of  
1,3,5-alkylsubstituted cyclohexylamines binding at  
NMDA receptor PCP site  
AUTHOR(S): Jirgensons, Aigars; Kauss, Valerjans; Kalvinsh, Ivars;  
Gold, Markus R.; Danysz, Wojciech; Parsons, Chris G.;  
Quack, Gunter  
CORPORATE SOURCE: Latvian institute of Organic Synthesis, Riga, LV-1006,  
Latvia  
SOURCE: European Journal of Medicinal Chemistry (2000), 35(6),  
555-565  
CODEN: EJMCA5; ISSN: 0223-5234  
PUBLISHER: Editions Scientifiques et Medicales Elsevier  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 133:281550  
REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 18 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2000:381463 CAPLUS  
DOCUMENT NUMBER: 133:17228  
TITLE: Preparation of polyalkylcyclohexane(alkan)amines as  
NMDA receptor antagonists  
INVENTOR(S): Gold, Markus; Danysz, Wojciech; Parsons, Christopher  
Graham Raphael; Kalvinsh, Ivars; Kauss, Valerjans;  
Jirgensons, Aigars  
PATENT ASSIGNEE(S): Merz & Co. Gmbh & Co., Germany  
SOURCE: U.S., 22 pp., Cont.-in-part of U.S. Ser. No. 48,575,  
abandoned.  
CODEN: USXXAM  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 3  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6071966	A	20000606	US 1998-141380	19980827
PRIORITY APPLN. INFO.:			US 1997-885944	B3 19970630
			US 1998-48575	B2 19980326

OTHER SOURCE(S): MARPAT 133:17228  
REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 19 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 2000:157723 CAPLUS  
DOCUMENT NUMBER: 132:194135  
TITLE: Preparation of cyclohexane(alkan)amines as drugs  
INVENTOR(S): Gold, Markus; Danysz, Wojciech; Parsons, Christopher  
Graham Raphael; Kalvinsh, Ivars; Kauss, Valerjans;  
Jirgensons, Aigars  
PATENT ASSIGNEE(S): Merz & Co. G.m.b.H. & Co., Germany  
SOURCE: U.S., 22 pp., Cont.-in-part of U.S. Ser. No. 885,944,  
abandoned.

CODEN: USXXAM  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 3  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6034134	A	20000307	US 1998-141381	19980827
PT 1009732	T	20031031	PT 1998-939579	19980624
CN 1136186	B	20040128	CN 1998-806775	19980624
ES 2200358	T3	20040301	ES 1998-939579	19980624
CZ 293248	B6	20040317	CZ 1999-4571	19980624
ZA 9805678	A	20000110	ZA 1998-5678	19980629
ZA 2002002908	A	20030714	ZA 2002-2908	20020412

PRIORITY APPLN. INFO.: US 1997-885944 B2 19970630  
OTHER SOURCE(S): MARPAT 132:194135  
REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 20 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 1999:159243 CAPLUS  
DOCUMENT NUMBER: 130:347295  
TITLE: Amino-alkyl-cyclohexanes are novel uncompetitive NMDA receptor antagonists with strong voltage-dependency and fast blocking kinetics: in vitro and in vivo characterization  
AUTHOR(S): Parsons, Chris G.; Danysz, Wojciech; Bartmann, Annette; Spielmanns, Peter; Frankiewicz, Tadeusz; Hesselink, Mayke; Eilbacher, Bernd; Quack, Gunter  
CORPORATE SOURCE: Department of Pharmacology, Merz + Co., Frankfurt am Main, D-60318, Germany  
SOURCE: Neuropharmacology (1999), 38(1), 85-108  
CODEN: NEPHBW; ISSN: 0028-3908  
PUBLISHER: Elsevier Science Ltd.  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
REFERENCE COUNT: 79 THERE ARE 79 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 21 OF 21 CAPLUS COPYRIGHT 2004 ACS on STN  
ACCESSION NUMBER: 1999:48692 CAPLUS  
DOCUMENT NUMBER: 130:119606  
TITLE: 1-amino-alkylcyclohexane NMDA receptor antagonists, preparation, and therapeutic use  
INVENTOR(S): Gold, Markus; Danysz, Wojciech; Parsons, Christopher  
Graham Raphael; Kalvinsh, Ivars; Kauss, Valerjans; Jirgensons, Aigars  
PATENT ASSIGNEE(S): Merz & Co. G.m.b.H. & Co., Germany  
SOURCE: PCT Int. Appl., 57 pp.  
CODEN: PIXXD2  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 3  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9901416	A2	19990114	WO 1998-EP4026	19980624
WO 9901416	A3	19990819		

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,

UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM  
 RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES,  
 FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI,  
 CM, GA, GN, ML, MR, NE, SN, TD, TG

AU 9888042	A1	19990125	AU 1998-88042	19980624
AU 724974	B2	20001005		
EP 1009732	A2	20000621	EP 1998-939579	19980624
EP 1009732	B1	20030521		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, LT, LV				
JP 2002511873	T2	20020416	JP 1999-506323	19980624
AT 240936	E	20030615	AT 1998-939579	19980624
PT 1009732	T	20031031	PT 1998-939579	19980624
CN 1136186	B	20040128	CN 1998-806775	19980624
ES 2200358	T3	20040301	ES 1998-939579	19980624
CZ 293248	B6	20040317	CZ 1999-4571	19980624
ZA 9805678	A	20000110	ZA 1998-5678	19980629
MX 9911993	A	20000930	MX 1999-11993	19991217
FI 9902801	A	19991229	FI 1999-2801	19991229
NO 9906548	A	20000228	NO 1999-6548	19991229
ZA 2002002908	A	20030714	ZA 2002-2908	20020412
PRIORITY APPLN. INFO.:			US 1997-885944	A 19970630
			WO 1998-EP4026	W 19980624
OTHER SOURCE(S):			MARPAT 130:119606	

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(FILE 'HOME' ENTERED AT 14:59:06 ON 25 JUN 2004)

FILE 'CAPLUS' ENTERED AT 14:59:23 ON 25 JUN 2004

L1 1 S US6071966/PN

SELECT L1 1 RN

L2 21 S E4-E120

FILE 'MEDLINE, BIOSIS, EMBASE' ENTERED AT 15:46:46 ON 25 JUN 2004

L3 189 S EMESIS(S) (CNS OR CENTRAL(3A)NERVOUS(4A)SYSTEM)

FILE 'MEDLINE' ENTERED AT 15:57:42 ON 25 JUN 2004

L4 14 S L3 NOT PY>=1999

L5 710 S (EMESIS OR VOMIT?) (L) (CNS OR CENTRAL(3A)NERVOUS(4A)SYSTEM)

L6 531 S L5 NOT PY>=1999

L7 6 S (APPETITE(4A)DISORDER#) (L) (CNS OR CENTRAL(3A)NERVOUS(4A)SYSTE

L8 12 S (CEREBELLAR(3A)TREMOR) (L) (CNS OR CENTRAL(3A)NERVOUS(4A)SYSTEM

=>

99597,102 ✓

ACCESSION NUMBER: 97081306 MEDLINE  
DOCUMENT NUMBER: PubMed ID: 9118822  
TITLE: Ondansetron. A review of its pharmacology and preliminary clinical findings in novel applications.  
AUTHOR: Wilde M I; Markham A  
CORPORATE SOURCE: Adis International Limited, Auckland, New Zealand.  
SOURCE: Drugs, (1996 Nov) 52 (5) 773-94. Ref: 185  
Journal code: 7600076. ISSN: 0012-6667.  
PUB. COUNTRY: New Zealand  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, ACADEMIC)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199704  
ENTRY DATE: Entered STN: 19970506  
Last Updated on STN: 19970506  
Entered Medline: 19970422

AB The use of ondansetron, a selective serotonin 5-HT<sub>3</sub> receptor antagonist, is well established in patients with nausea and vomiting associated with cancer chemotherapy, radiotherapy or anaesthesia and surgery. The wide distribution of 5-HT<sub>3</sub> receptors in the body and the role of these receptors in disease have provided the rationale for investigation of ondansetron in novel applications. Preliminary data have shown ondansetron to have clinical benefit in patients with nausea and vomiting associated with drug overdosage or poisoning, anti-infective or antidepressant therapies, uraemia or neurological trauma, and in patients with pruritus. Patients with gastrointestinal motility disorders (e.g. carcinoid syndrome, irritable bowel syndrome, diarrhoea associated with cryptosporidiosis or diabetes, and chronic refractory diarrhoea) have also shown some improvement when treated with ondansetron, as have patients with certain pain or CNS-related disorders [e.g. alcohol (ethanol) dependence, opiate withdrawal, vertigo, cerebellar tremor and Parkinson's disease treatment-related psychosis]. In contrast to conventional antiemetics, ondansetron is generally well tolerated with a lower incidence of sedation and only isolated case reports of extrapyramidal reactions. Furthermore, unlike dopamine receptor-blocking neuroleptics, ondansetron does not appear to worsen the symptoms of Parkinson's disease. Thus, in addition to its established indications, preliminary results suggest that ondansetron may be beneficial in a number of novel applications. This drug may represent a treatment alternative in patients with refractory disease, or an effective treatment of conditions for which current therapies are either poorly tolerated or not available. Further investigation of ondansetron in a range of potential new applications appears to be warranted.

87/597, 102

ACCESSION NUMBER: 1998105332 MEDLINE  
DOCUMENT NUMBER: PubMed ID: 9443121  
TITLE: Vomiting and gastroesophageal motor activity in  
children with disorders of the central  
nervous system.  
COMMENT: Comment in: J Pediatr Gastroenterol Nutr. 1998  
Sep;27(3):373-4. PubMed ID: 9740220  
AUTHOR: Ravelli A M; Milla P J  
CORPORATE SOURCE: Department of Gastroenterology, Institute of Child Health,  
London, United Kingdom.  
SOURCE: Journal of pediatric gastroenterology and nutrition, (1998  
Jan) 26 (1) 56-63.  
Journal code: 8211545. ISSN: 0277-2116.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 199802  
ENTRY DATE: Entered STN: 19980226  
Last Updated on STN: 20000303  
Entered Medline: 19980213

AB BACKGROUND: Vomiting is common in children with disorders of the  
central nervous system (CNS) and is  
usually ascribed to gastroesophageal reflux (GER). However, recent  
acquisitions on the pathophysiology of vomiting suggest that the  
dysmotility of the foregut may be more widespread. METHODS: Fifty-five  
children with CNS disorders, 50 of whom suffered from retching  
and/or vomiting (18 following fundoplication) were studied. We  
assessed GER by 24 hour pH monitoring and endoscopy, gastric electrical  
activity by electrogastrography, and gastric half-emptying time (T1/2) of  
a milk meal by electrical impedance tomography. RESULTS: Of the 50  
vomiting patients, 29 had GER (reflux index of 5.7%-87.4%;  
controls: < 5%), and 31 had gastric dysrhythmias (12 tachyarrhythmia at  
5.5-11.2 cpm, 4 bradyarrhythmia at 1.7-1.9 cpm, 15 unstable electrical  
activity; controls; 2.2-4.0 cpm). Sixteen patients had GER and gastric  
dysrhythmias. Eleven of 18 patients with fundoplication had gastric  
dysrhythmias. Gastric T1/2 was delayed in 12 of 13 patients with gastric  
dysrhythmia (6 with GER), versus 2 of 5 with GER alone. No abnormalities  
were detected in the 5 patients who did not suffer from vomiting  
. CONCLUSIONS: Children with CNS disorders who vomit  
have abnormal gastric motility as often as GER. Following fundoplication,  
many patients continue to have symptoms possibly related to gastric  
dysrhythmias, the effects of which may be unmasked by fundoplication.  
Foregut dysmotility may be related to abnormal modulation of the enteric  
nervous system by the CNS or to involvement of the enteric  
nervous system by the same process affecting the brain.

09/597, 62

ACCESSION NUMBER: 85102106 MEDLINE  
DOCUMENT NUMBER: PubMed ID: 2857138  
TITLE: Mechanisms of appetite modulation by drugs.  
AUTHOR: Sullivan A C; Gruen R K  
SOURCE: Federation proceedings, (1985 Jan) 44 (1 Pt 1) 139-44.  
Ref: 51  
Journal code: 0372771. ISSN: 0014-9446.  
PUB. COUNTRY: United States  
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
LANGUAGE: English  
FILE SEGMENT: Priority Journals  
ENTRY MONTH: 198502  
ENTRY DATE: Entered STN: 19900320  
Last Updated on STN: 19970203  
Entered Medline: 19850225

AB The regulation of appetite is a complex process that we are just beginning to understand. It consists of both central and peripheral elements and involves the integration by the brain of a variety of signals from peripheral organs transmitted by neurotransmitters, peptides, hormones, and metabolites. All available anorectic drugs act by central mechanisms and have several disadvantages including limited effectiveness, side effects on the **central nervous system**, the development of tolerance, abuse potential, and rebound hyperphagia on discontinuance. Several appetite-modulating agents have been tested in animals that act by peripheral mechanisms and do not produce tolerance or rebound hyperphagia, which suggests that peripherally acting anorectic drugs may provide novel therapeutic approaches to **disorders of appetite** regulation in humans.